

EC DECLARATION OF CONFORMITY

RF19-0089 Rev. D
ECN Number: 22-03480

We, MicroVention Europe SARL, located in Saint-Germain-en-Laye, France, declare according to Directive 93/42/EEC Annex II under our sole responsibility that the products to which this declaration relates are in conformity with Directive 93/42/EEC and fulfill the Essential Requirements as described in Directive 93/42/EEC Annex I.

Council Directive 93/42/EEC

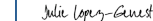
Conformity Assessment Procedure Performed:

EC Design Examination Certificate <input checked="" type="checkbox"/> <u>(Annex II.4)</u> 517356 MRA Certificate Number	EC Full Quality Assurance Certificate <input checked="" type="checkbox"/> <u>(Annex II.3)</u> 487703 MR2 Certificate Number
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Product	Model Number(s)	Class/Rule	GMDN Code
PHIL™ Liquid Embolic System	LEN10250 LEN10250RE LEN10300 LEN10300RE LEN10350 LEN10350RE LEN10LV250 LEN10LV250RE	Class III, Annex 9, Rule 8	60939

Legal Manufacturer	Production Site(s)	Notified Body
MicroVention Europe SARL 30 bis, rue du Vieil Abreuvoir 78100 Saint-Germain-en-Laye France	MicroVention Inc. 35 Enterprise Aliso Viejo, CA 92656 USA	DQS Medizinprodukte GmbH D-60433 Frankfurt am Main, Germany Notified Body No: 0297

We herewith declare that the above-mentioned medical device (s) meet the provisions of the council directive 93/42/EEC Medical Device Directive. This declaration is supported by the EC Quality System Certificate (s) according to the provisions of the relevant Annex (es) of above Directive. This declaration applies to all device(s) specified above distributed from the signature date forward.

DocuSigned by:

 Signer Name: Julie Lopez-Genest
 Signing Reason: I approve this document
 Signing Time: 9/9/2022 | 9:17:14 AM PDT
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Saint-Germain-en-Laye, France Place of Issue	9/9/2022 Date of Issue
Julie Lopez-Genest Sr Manager, RA & QA, EMEA MicroVention Inc.	

Certificate Expiry Date: 2024-05-26